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NOTICE OF ALLOWANCE AND FEE(S) DUE

30565 7590 12/14/2009

12/14/2009

WOODARD, EMHARDT, MORIARTY, MCNETT & HENRY LLP 111 MONUMENT CIRCLE, SUITE 3700 INDIANAPOLIS. IN 46204-5137 EXAMINER

GUPTA, ANISH

ART UNIT PAPER NUMBER

1654 DATE MAILED: 12/14/2009

 APPLICATION NO.
 FILING DATE
 FIRST NAMED INVENTOR
 ATTORNEY DOCKET NO.
 CONFRMATION NO.

 10/551,840
 01/20/2006
 Fabrizio Samaritani
 7541-6
 4228

TITLE OF INVENTION: LIQUID PHARMACEUTICAL FORMULATIONS OF FSH AND LH TOGETHER WITH A NON-IONIC SURFACTANT

 APPLN. TYPE
 SMALL ENTITY
 ISSUE FEE DUE
 PUBLICATION FEE DUE
 PREV. PAID ISSUE FEE
 TOTAL FEES) DUE
 DATE DUE

 nonprovisional
 NO
 \$1510
 \$300
 \$0
 \$1810
 03/15/2010

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. PROSECUTION ON THE MERITS IS CLOSED. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN THREE MONTHS FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. THIS STATUTORY PERIOD CANNOT BE EXTENDED. SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

HOW TO REPLY TO THIS NOTICE:

I. Review the SMALL ENTITY status shown above.

If the SMALL ENTITY is shown as YES, verify your current SMALL ENTITY status:

A. If the status is the same, pay the TOTAL FEE(S) DUE shown above.

B. If the status above is to be removed, check box 5b on Part B Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and twice the amount of the ISSUE FEE shown above, or

If the SMALL ENTITY is shown as NO:

A. Pay TOTAL FEE(S) DUE shown above, or

B. If applicant claimed SMALL ENTITY status before, or is now claiming SMALL ENTITY status, check box 5a on Part B - Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and I/2 the ISSUE FIEE shown above.

II. PART B - FEE(S) TRANSMITTAL, or its equivalent, must be completed and returned to the United States Patent and Trademark Office (USPTO) with your ISSUE FEE and PUBLICATION FEE (if required). If you are charging the fee(s) to your deposit account, section "4b" of Part B - Fee(s) Transmittal should be completed and an extra copy of the form should be submitted. If an equivalent of Part B is filed, a request to reapply a previously paid issue fee must be clearly made, and delays in processing may occur due to the difficulty in recognizing the paper as an equivalent of Part B.

III. All communications regarding this application must give the application number. Please direct all communications prior to issuance to Mail Stop ISSUE FEE unless advised to the contrary.

IMPORTANT REMINDER: Utility patents issuing on applications filed on or after Dec. 12, 1980 may require payment of maintenance fees. It is patentee's responsibility to ensure timely payment of maintenance fees when due.

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Complete and send this form, together with applicable fee(s), to: Mail Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450

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appropriate. All further indicated unless correct maintenance fee notifica	ed belôw or directed oth	ng the Patent, advance on herwise in Block 1, by (rders and notification a) specifying a new co	of ma	aintenance fees w ondence address;	ill be and/or	mailed to the current (b) indicating a sepa	correspond rate "FEE	Jence address as ADDRESS" for
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10/551,840			Fabrizio Samaritani		7541-6 4228			4228	
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APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE D	UE I	PREV. PAID ISSUE	SPEE	TOTAL FEE(S) DUE		ATE DUE
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(A) NAME OF ASSI	less an assignee is ident h in 37 CFR 3.11. Comp GNEE	ified below, no assignee pletion of this form is NO	data will appear on the transmission of tr	he pateg an as	ent. If an assign ssignment. and STATE OR C	OUNT	'RY)		
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4a. The following fee(s) Issue Fee Publication Fee (N		4b, Payment of Fee(s): (Please first reapply any previously paid issue fee shown above) A check is enclosed. Payment by credit card. Form PTO-2038 is attached. The Director is hereby authorized to charge the required fee(s), any deficiency, or credit any overpayment, to Deposit Account Number (enclose an extra copy of this form).							
	s SMALL ENTITY state	as. See 37 CFR 1.27.					ITTY status. Sec 37 Cl		
NOTE: The Issue Fee an interest as shown by the	d Publication Fee (if req records of the United Sta	uired) will not be accepte tes Patent and Trademark	d from anyone other the Office.	nan the	e applicant; a regi	stered a	attorney or agent; or th	e assignee	or other party in
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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
10/551,840	01/20/2006	Fabrizio Samaritani	7541-6	4228		
30565	7590 12/14/2009	EXAMINER				
WOODARD, E	MHARDT, MORIAR	GUPTA, ANISH				
	T CIRCLE, SUITE 370	ART UNIT	PAPER NUMBER			
INDIANAPOLIS	, IN 46204-5137	1654				

DATE MAILED: 12/14/2009

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment to date is 0 day(s). If the issue fee is paid on the date that is three months after the mailing date of this notice and the patent issues on the Tuesday before the date that is 28 weeks (six and a half months) after the mailing date of this notice, the Patent Term Adjustment will be 0 day(s).

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

Application No. Applicant(s) 10/551.840 SAMARITANI ET AL. Notice of Allowability Examiner Art Unit ANISH GUPTA 1654 -- The MAILING DATE of this communication appears on the cover sheet with the correspondence address--All claims being allowable, PROSECUTION ON THE MERITS IS (OR REMAINS) CLOSED in this application. If not included herewith (or previously mailed), a Notice of Allowance (PTOL-85) or other appropriate communication will be mailed in due course. THIS NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHTS. This application is subject to withdrawal from issue at the initiative of the Office or upon petition by the applicant. See 37 CFR 1.313 and MPEP 1308. This communication is responsive to 7-06-09. The allowed claim(s) is/are 46-49,51-53,57-63,72-80,82-89,189-193,195-201 and 203-217. 3. Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). b) ☐ Some* c) ☐ None of the: a) 🔯 All 1. A Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this national stage application from the International Bureau (PCT Rule 17.2(a)). * Certified copies not received: _____. Applicant has THREE MONTHS FROM THE "MAILING DATE" of this communication to file a reply complying with the requirements noted below. Failure to timely comply will result in ABANDONMENT of this application. THIS THREE-MONTH PERIOD IS NOT EXTENDABLE. A SUBSTITUTE OATH OR DECLARATION must be submitted. Note the attached EXAMINER'S AMENDMENT or NOTICE OF INFORMAL PATENT APPLICATION (PTO-152) which gives reason(s) why the oath or declaration is deficient. CORRECTED DRAWINGS (as "replacement sheets") must be submitted. (a) Including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached 1) hereto or 2) to Paper No./Mail Date (b) including changes required by the attached Examiner's Amendment / Comment or in the Office action of Identifying indicia such as the application number (see 37 CFR 1.84(c)) should be written on the drawings in the front (not the back) of each sheet. Replacement sheet(s) should be labeled as such in the header according to 37 CFR 1.121(d). 6. DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL. Attachment(s) 1. | Notice of References Cited (PTO-892) 5. Notice of Informal Patent Application 2. Notice of Draftperson's Patent Drawing Review (PTO-948) Interview Summary (PTO-413), Paper No./Mail Date Information Disclosure Statements (PTO/SB/08). 7. X Examiner's Amendment/Comment Paper No./Mail Date 4. T Examiner's Comment Regarding Requirement for Deposit 8. X Examiner's Statement of Reasons for Allowance of Biological Material 9. ☐ Other . /Anish Gupta/

Primary Examiner, Art Unit 1654

DETAILED ACTION

EXAMINER'S AMENDMENT

An examiner's amendment to the record appears below. Should the changes and/or
additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312.
 To ensure consideration of such an amendment, it MUST be submitted no later than the payment of
the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Thomas Henry on September 24, 2009.

The application has been amended as follows:

The following claims have been amended as follows:

- 46. A liquid pharmaceutical composition, comprising:
 - a follicle stimulating hormone or a variant thereof,
 - m-cresol.
 - a diluent, and
 - poloxamer 188,

the composition having a stability sufficient to avoid precipitation.

- A liquid pharmaceutical composition, comprising:
 - a follicle stimulating hormone or a variant thereof,
 - a luteinising hormone or a variant thereof,
 - a bacteriostatic agent selected form the group consisting of phenol and m-cresol,
 - poloxamer 188, and
 - a diluent

wherein <u>either</u> the follicle stimulating hormone is human follicle stimulating hormone, <u>or</u> the luteinising hormone is human luteinising hormone, or <u>both</u> the follicle stimulating hormone is human follicle stimulating hormone and the luteinising hormone is human luteinising hormone,

the composition having a stability sufficient to avoid precipitation.

73. A liquid pharmaceutical composition, comprising:

- a follicle stimulating hormone or a variant thereof,
- a luteinising hormone or a variant thereof,
- a bacteriostatic agent selected form the group consisting of phenol and m-cresol, poloxamer 188, and
- a diluent

wherein either the follicle stimulating hormone is urinary human follicle stimulating hormone, or the luteinising hormone is urinary human luteinising hormone, or both the follicle stimulating hormone is urinary human follicle stimulating hormone and the luteinising hormone is urinary human luteinising hormone,

the composition having a stability sufficient to avoid precipitation.

74. A liquid pharmaceutical composition, comprising:

- a follicle stimulating hormone or a variant thereof,
- a luteinising hormone or a variant thereof,
- a bacteriostatic agent selected form the group consisting of phenol and m-cresol, poloxamer 188, and

a diluent

wherein <u>either</u> the follicle stimulating hormone is recombinant human follicle stimulating hormone, <u>or</u> the luteinising hormone is recombinant human luteinising hormone, or both the follicle stimulating hormone is recombinant human follicle stimulating hormone and the recombinant luteinising hormone is human luteinising hormone,

198. A liquid pharmaceutical composition, comprising:

a follicle stimulating hormone or a variant thereof.

phenol,

a diluent, and

poloxamer 188,

the composition having a stability sufficient to avoid precipitation.

the composition having a stability sufficient to avoid precipitation.

The following is an examiner's statement of reasons for allowance:

The claims are drawn to a liquid composition comprising follicle stimulating hormone and poloxamer 188 as two of the specific components.

Hoffman et al. teaches a formulation comprising follicle stimulating hormone. The reference states "Other additives, such as a pharmaceutically acceptable solubilizers like Tween 20 (polyoxyethylene (20) sorbitan monopalmitate), Tween 40 (polyoxyethylene (20) sorbitan monopalmitate), Tween 80 (polyoxyethylene (20) sorbitan monopalmitate), Pluronic F68 (polyoxyethylene polyoxypropylene block copolymers), and PEG (polyothylene glycol) or non-ionic

surfactants such as polysorbate 20 or 80 or poloxamer 184 or 188, Pluronic.RTM. polyls, other block co-polymers, and chelators such as EDTA and EGTA may optionally be added to the formulations or compositions to reduce aggregation." (see paragraph [0100]). Thus, reading the reference, one would expect both poloxamer and tween 20 to behave similarly.

However, as Applicants specification asserts formulations with FSH and mixtures of FSH and LH with a Pluronic F68 (BASF, Pluronic F68 is also known as Poloxamer 188) obtain a stable formulation that avoids the problem of precipitation in the presence of a bacteriostatic agent, such as m-cresol and phenol. Precipitation, resulting in the formation of turbid or milky solutions occurs when TWEEN 20 is used with m-cresol or phenol (see page 12 of the specification). This is unexpected since one would have expected to observe the same "turbid or milky solution" for poloxamer 188 based on tween 20. Based on Hoffman et al.'s disclosure one would predict Tween 20 and poloxamer 188 to behave the same way. Furthermore, Hoffman et al. does not teach nor suggest the use of Poloxamer 188 over Tween 20, PEG or other surfactant listed. Indeed, Hoffman does not provide a single example that uses either Tween 20 or Poloxamer. Thus, based on Applicants disclosure that formulations with FSH and mixtures of FSH and LH with a Pluronic F68 (BASF, Pluronic F68 is also known as Poloxamer 188) obtain a stable formulation that avoids the problem of precipitation in the presence of a bacteriostatic agent, such as m-cresol and phenol, and the fact that Hoffman does not provide any specific motivation to choose Poloxamer 188 from the list of additives, the claims of the instant application are both novel and unobvious.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

2. Any inquiry concerning this communication or earlier communications from the examiner should be directed to ANISH GUPTA whose telephone number is (571)272-0965. The examiner can normally be reached on 5/4/9.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Tsang Cecilia can be reached on 571-272-0562. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Anish Gupta/ Primary Examiner, Art Unit 1654